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(51) International Patent Classification ⁶ : A61K 9/16	A2	(11) International Publication Number: WO 97/10810 (43) International Publication Date: 27 March 1997 (27.03.97)
(21) International Application Number: PCT/JP96/02504 (22) International Filing Date: 4 September 1996 (04.09.96) (30) Priority Data: 9518465.1 9 September 1995 (09.09.95) GB (71) Applicant (for all designated States except US): SMITHKLINE BEECHAM SEIYAKU KK [JP/JP]; SB Building, 6, Sanban-cho, Chiyoda-ku, Tokyo 102 (JP). (72) Inventors; and (75) Inventors/Applicants (for US only): AKIYAMA, Hidero [JP/JP]; SmithKline Beecham Seiyaku KK, Takasaki Laboratories, 168, Ohyagi-machi, Takasaki-shi, Gunma 370 (JP). MATSUMOTO, Zene [JP/JP]; SmithKline Beecham Seiyaku KK, Takasaki Laboratories, 168, Ohyagi-machi, Takasaki-shi, Gunma 370 (JP). UENO, Takashi [JP/JP]; SmithKline Beecham Seiyaku KK, Takasaki Laboratories, 168, Ohyagi-machi, Takasaki-shi, Gunma 370 (JP). (74) Agents: AOYAMA, Tamotsu et al.; Aoyama & Partners, IMP Building, 3-7, Shiromi 1-chome, Chuo-ku, Osaka-shi, Osaka 540 (JP).		(81) Designated States: AL, AM, AU, AZ, BB, BG, BR, BY, CA, CN, CU, CZ, EE, GE, HU, IL, IS, JP, KG, KR, KZ, LC, LK, LR, LT, LV, MD, MG, MK, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, TJ, TM, TR, TT, UA, US, UZ, VN, ARIPO patent (KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>Without international search report and to be republished upon receipt of that report.</i>
(54) Title: PHARMACEUTICAL GRANULES (57) Abstract. A high speed agitation granulator method of preparing a substantially spherical granule for pharmaceutical use comprising a medicament for pharmaceutical use, wherein the medicament has an aqueous solubility of 0.01 to 0.30 g/ml, which method comprises introducing a mixture of medicament and excipients composing at least 5 % crystalline cellulose into the granulator and spraying on water or a mixture of ethanol and water as binder solution; a substantially spherical granule for pharmaceutical use comprising famciclovir and 5 % or more crystalline cellulose, together with an optional coating; and a sachet containing a unit dose of famciclovir in the form of such granules.		

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DESCRIPTION

PHARMACEUTICAL GRANULES

FIELD OF THE INVENTION

This invention relates to a method of preparation
5 of granules incorporating a medicament for pharmaceutical use.

BACKGROUND OF THE INVENTION

Granules are a suitable form for medicaments which
are to be administered orally. A dose of medicament in the
form of granules may be measured and administered by
10 incorporating into a sachet or using a spoon or may be
incorporated into a capsule or formulated into a tablet to be
swallowed. It is preferred that such granules are of regular
shape and substantially spherical to give good fluidity for
ease of dispensing and capsule filling or tableting as well
15 as for aesthetic appeal. Spherical granules are also easier
to coat with taste masking, enteric/protective and sustained
release coating materials.

At present, there are two methods of preparing
spherical granules. One is to extrude a cylindrical granule
20 using a "squeeze-out" type granulation machine and then to
make it spherical in shape by using a marumeriser, a machine
which "cuts off the edges" of the cylinder shape. This method

involves difficult manufacturing conditions and is time consuming which precludes its routine use. The other method used is to gradually coat a core particle such as granulated sugar using a centrifugal flow type granulator but this is a time consuming process and is not generally applicable to a wide range of medicaments where the proportion of the volume of the medicament in the granules high (20% or higher).

The most efficient method of preparing granules is using a high-speed agitation granulator machine which mixes and granulates in one operation using two rotating blades, a main blade with a horizontal plane of rotation and a chopper blade above it with a vertical plane of rotation (cross blade). The medicament and excipients are introduced into the machine in advance and binder solution is then poured or sprayed into the machine from above whilst the blades are rotating. By this method, granules can be prepared quickly and easily but the disadvantage is that the resulting granules are irregular and non-spherical in shape and do not have the advantages associated with spherical granules.

Surprisingly, we have now discovered a method of preparing substantially spherical granules using a high-speed agitation granulator machine thus avoiding the need to use the "squeeze-out" and marumerizer method.

DETAILED DESCRIPTION OF THE INVENTION

Accordingly, the present invention provides a high speed agitation granulator method of preparing a substantially spherical granule for pharmaceutical use comprising a medicament for pharmaceutical use wherein the medicament has an aqueous solubility of 0.01 to 0.30 g/ml, which method comprises introducing a mixture of the medicament and excipients comprising at least 5% crystalline cellulose into the granulator and spraying on water or a mixture of ethanol and water as binder solution.

Suitable medicaments include caffeine which has an aqueous solubility 0.02 g/ml, pyridoxine hydrochloride which has an aqueous solubility of 0.22 g/ml and particularly the orally administered antiviral compounds, famciclovir and acyclovir which have a aqueous solubility of more than 0.25 g/ml. Generally the proportion of medicament with respect to excipients in the composition of the granule is up to 25%, such as up to 5%, 10%, 20%, although it is envisaged that up to 55% medicament could be incorporated.

The mixture of medicament and excipients comprise at least 5% crystalline cellulose, for example up to 10%, 20%, 30%, 40%, 50%, 60%, 70%, 80% and 90%. Other suitable additional excipients include lactose, although the highest yield of spherical granules are when the mixture contains crystalline cellulose as the sole excipient, for example where

the composition contains 25% medicament and 75% crystalline cellulose.

The binder solution may be up to 75% ethanol and it is found that the sphericity of granules is higher when the ethanol concentration is higher although from a manufacturing environment point of view and to avoid any problems resulting from residual organic solvent, it may be preferable to use pure water as the binder solution. A linear relationship between the granule size and the amount of binder solution was observed such that the granule size can be regulated by this method (the greater the amount of binder solution, the larger the granule). The granules suitable for administration by spoon are generally more than 500 μm in diameter, favourably more than 700 μm , up to 1500 μm , favourably up to 1000 μm . Granules suitable for capsules or tablets are usually smaller than 500 μm .

The granules are suitably agitated in the granulator after spraying of the binder solution to improve the spherical shape and smooth the surface of the granules. The agitation time will depend on the size of the granules and composition of the granules, together with the size of the granulator and the speed of rotation of the main blade and the cross blade. The agitation times generally range from 2 to 30 minutes.

The invention provides, in one aspect, a substantially spherical granule for pharmaceutical use

comprising famciclovir and 5% or more crystalline cellulose, together with an optional coating. In a preferred aspect the invention provides such a granule comprising famciclovir and crystalline cellulose as the sole excipient, for example a granule containing 25% famciclovir and 75% cellulose.

The following Examples illustrate the process of the invention. The accompanying Figures and Tables illustrate the following with respect to pyridoxine granules:

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 shows photographs of granules taken through a microscope (x5). (A) is where the binder solution is water and (B) is where the binder solution is 70% ethanol.

Figure 2 shows photographs of granules taken through a microscope (x5). (A) is with 20% crystalline cellulose and (B) is with 75% crystalline cellulose.

Figure 3 is a graph showing the relationship between amount of binder solution and the size of granules.

Figure 4 is a graph showing the relationship between agitation time of the main and cross blades with respect to the shape factor of granules.

Figure 5 (A), (B-1), and (C) show photographs of granules taken through a microscope (x5) where agitation times are 0, 14 and 30 minutes respectively. Figure 5 (B-2) shows

a photograph of granules taken through a microscope (15) where the agitation time is 14 minutes.

Figure 6 is a graph shows the dissolution curve of the granules.

5 Figure 7 shows a photograph of granules taken through a microscope (x10) after the dissolution test.

Table I shows the various compositions of the granules with respect to crystalline cellulose content.

10 Table II shows the effect of ethanol concentration in the binder solution on Shape Factor and yield of granules.

Table III shows the effect of microcrystalline cellulose concentration on Shape Factor and yield of granules.

15 Table IV shows the effect of amount of binder solution on Particle Size Distribution of 25% pyridoxine granules.

Table V shows the reproducibility of Shape Factor of 25% pyridoxine granules.

Table VI shows the reproducibility of Particle Size Distribution and yield of 25% pyridoxine granules.

EXAMPLESPreparation Example 1 (10% caffeine spherical granule)

In this example, 700 g of a mixture having the composition shown below were first charged into an agitation granulator (Type VG-5, manufactured by Powlex Corp., Japan) and then 180 ml of 20% ethanol-water solution as a binder liquid were sprayed, at a rate of 100 ml/min, over the mixture in the granulator while rotating the main blade and the cross blade of the granulator at 600 and 3000 rpm, respectively. After the end of spraying, the granulating operation was continued for another two minutes. The resulting grain was then dried at 50°C for 12 hours using a convection drying oven to obtain the spherical granule.

Composition of the mixture:

Caffeine	10 parts
Lactose	82 parts
Crystalline cellulose	8 parts

The particle size of this spherical granule obtained was 500 - 850 um with a yield of 85.5%.

Preparation Example 2 (25% caffeine spherical granule)

In this example, 700 g of a mixture having the composition shown below were first charged into an agitation granulator (Type VG-5, manufactured by Powlex Corp., Japan)

and then 370 ml of 70% ethanol-water solution as a binder liquid were sprayed, at a rate of 100 ml/min, over the mixture in the granulator while rotating the main blade and the cross blade of the granulator at 600 and 3000 rpm, respectively. After the end of spraying, the granulating operation was continued for another five minutes while rotating the main blade and the cross blade at 300 and 1000 rpm, respectively. The resulting grain was then dried at 50°C for 12 hours using a convection drying oven to obtain the spherical granule.

Composition of the mixture:

Caffeine	25 parts
Lactose	55 parts
Crystalline cellulose	20 parts

The particle size of this spherical granule obtained was 710 - 1180 um with a yield of 77.7%.

Preparation Example 3 (25% pyridoxine hydrochloride spherical granule)

In this example, 500 g of a mixture having the composition shown below were first charged into an agitation granulator (Type VG-5, manufactured by Powlex Corp., Japan) and then 350 ml of water as a binder liquid were sprayed, at a rate of 100 ml/min, over the mixture in the granulator while rotating the main blade and the cross blade of the granulator at 600 and 3000 rpm, respectively. After completing the spraying, the granulating operation was continued for another

14 minutes while rotating the main blade and the cross blade at 300 and 1000 rpm, respectively. The resulting grain was then dried at 50°C for 12 hours using a convection drying oven to obtain the spherical granule.

5 Composition of the mixture:

Pyridoxine hydrochloride	25 parts
Crystalline cellulose	75 parts

The particle size of this spherical granule obtained was 710 - 1180 um with a yield of 96.4%.

10 Preparation Example 4 (25% famciclovir spherical granule)

In this example, 500 g of a mixture having the composition shown below were first charged into an agitation granulator (Type VG-5, manufactured by Powlex Corp., Japan) and then 458 or 459 ml of water as a binder liquid were sprayed, at a rate of 65 ml/min, over the mixture in the granulator while rotating the main blade and the cross blade of the granulator at 600 and 3000 rpm, respectively. After completing the spraying, the granulating operation was continued for another 30 minutes while rotating the main blade and the cross blade at 300 and 1000 rpm, respectively. The resulting grain was then dried at 50°C for 12 hours using a convection drying oven to obtain the spherical granule.

Composition of the mixture:

Famciclovir	25 parts
Crystalline cellulose	75 parts

Using 458 ml of binder solution, the particle size of the spherical granule obtained was 600 - 850 μ m with a yield of 97.1 percent. Using 459 ml of binder solution, the particle size of the spherical granule obtained was 710 - 1000 μ m with a yield of 97.0 percent.

Preparation Example 5 (25% pyridoxine hydrochloride spherical granule)

In this example, 500 g of a mixture having the composition shown below were first charged into an agitation granulator (Type VG-5, manufactured by Powlex Corp., Japan) and then 326 ml of water as a binder liquid were sprayed, at a rate of 80 ml/min, over the mixture in the granulator while rotating the main blade and the cross blade of the granulator at 600 and 3000 rpm, respectively. After completing the spraying, the granulating operation was continued for another 30 minutes while rotating the main blade and the cross blade at 400 and 2000 rpm, respectively. The resulting grain was then dried at 50°C for 12 hours using a convection drying oven to obtain the spherical granule.

Composition of the mixture:

Pyridoxine hydrochloride	25 parts
Crystalline cellulose	75 parts

The particle size of this spherical granule obtained was 300 - 500 um with a yield of 90.7%.

TABLE I. Formulas of Pyridoxine Spherical Granules(%)

Formulation No.	PSG-1	PSG-2	PSG-3	PSG-4	PSG-5	PSG-6	PSG-7	PSG-8
Pyridoxine hydrochloride	25	25	25	25	25	25	25	25
Microcrystalline cellulose	10	20	30	40	50	60	70	75
Lactose	65	55	45	35	25	15	5	0

15

TABLE II. Effect of Ethanol Concentration in the binder solution on Shape Factor and Yield of Pyridoxine Spherical Granules

	Ethanol concentration (%)	Shape Factor		Yield (%)
		SF1	SF2	
	0	124.4	111.1	88.9
	10	120.3	111.3	85.3
25	30	119.5	110.3	85.6
	50	118.6	109.7	89.4
	70	117.1	109.0	85.0

TABLE III. Effect of microcrystalline Cellulose Concentration on Shape Factor and Yield of Pyridoxine Spherical Granules

	Concentration of microcrystalline cellulose (%)	Shape Factor		Yield (%)
		SF1	SF2	
10	10	120.6	111.1	70.3
	20	119.5	109.8	79.3
	30	119.7	110.1	83.7
	40	119.4	109.6	84.5
	50	119.1	111.8	88.9
	60	117.3	109.4	94.3
	70	115.9	109.9	95.8
	75	115.6	109.5	96.6

TABLE IV. Effect of Amount of Binder Solution on Particle Size Distribution of 25% Pyridoxine Spherical Granules

	Particle size (μ m)	Amount of binder solution (ml)			
		490	500	510	520
20	1400 - 1700	0.3%	0.5%	0.9%	1.5%
	1180 - 1400	0.5	1.5	49.0	82.4
25	1000 - 1180	1.1	11.9	42.5	12.3
	850 - 1000	1.9	74.0	5.9	2.5
	710 - 850	7.5	10.6	1.1	0.6
	500 - 710	87.7	1.3	0.4	0.3
	355 - 500	0.8	0.1	0.1	0.1
30	0 - 355	0.2	0.1	0.1	0.3
D50%		617 μ m	875 μ m	1180 μ m	1270 μ m

TABLE V. Reproducibility of Shape Factor of 25% Pyridoxine Spherical Granules

	Repetition No.	SF1	SF2
5	1	114.1	108.2
	2	114.3	108.0
	3	114.2	107.9
	4	114.4	108.2
10			

TABLE VI. Reproducibility of Particle Size Distribution and Yield of 25% Pyridoxine Spherical Granules

	Particle size (μ m)	Repetition No.			
		No. 1	No. 2	No. 3	No. 4
15	1400 - 1700	0.9	0.2	0.2	0.4
	1180 - 1400	1.8	1.0	1.0	1.6
20	1000 - 1180	5.8	4.1	3.4	3.5
	850 - 1000	61.7	67.4	61.5	66.3
	710 - 850	27.7	24.9	30.8	25.7
	500 - 710	2.1	2.1	2.9	2.1
	355 - 500	0.0	0.2	0.1	0.4
25	0 - 355	0.0	0.0	0.1	0.0
	Yield (710 - 1180)	95.2	96.4	95.7	95.5

CLAIMS

1. A high speed agitation granulator method of preparing a substantially spherical granule for pharmaceutical use comprising a medicament for pharmaceutical use, wherein the
5 medicament has an aqueous solubility of 0.01 to 0.30 g/ml, which method comprises introducing a mixture of medicament and excipients comprising at least 5% crystalline cellulose into the granulator and spraying on water or a mixture of ethanol and water as binder solution.

10 2. A method according to claim 1, wherein the medicament is caffeine which has an aqueous solubility 0.02 g/ml, pyridoxine hydrochloride which has an aqueous solubility of 0.22 g/ml or famciclovir and acyclovir which have an aqueous solubility of more than 0.25 g/ml.

15 3. A method according to claim 2 wherein the medicament is famciclovir.

4. A method according to claim 1, 2 or 3 wherein the proportion of medicament with respect to excipients in the composition of the granule is up to 25%.

20 5. A method according to any one of claims 2 to 4 wherein the mixture of medicament and excipients comprise at least 5% crystalline cellulose as the sole excipient.

6. A method according to any one of claims 1 to 5 where the amount of binder solution is regulated such that

granules are suitable for administration by spoon and are generally more than 500 um in diameter.

7. A method according to any one of claims 1 to 5 wherein the granules are agitated in the granulator after spraying of the binder solution to improve the spherical shape and smooth the surface of the granules.

8. A method according to claim 1, substantially as described herein with reference to any one of the Examples.

9. A substantially spherical granule for pharmaceutical use comprising famciclovir and 5% or more crystalline cellulose, together with an optional coating.

10. A granule according to claim 9 comprising famciclovir and crystalline cellulose as the sole excipient.

11. A granule according to claim 10 containing approximately 25% famciclovir and approximately 75% cellulose.

12. A granule according to claim 9, substantially as described herein with reference to any one of the Examples.

13. A granule according to claim 11 having a diameter of 500 to 1500 um.

14. A sachet containing a unit dose of famciclovir in the form of granules as defined in any one of claims 9 to 13.

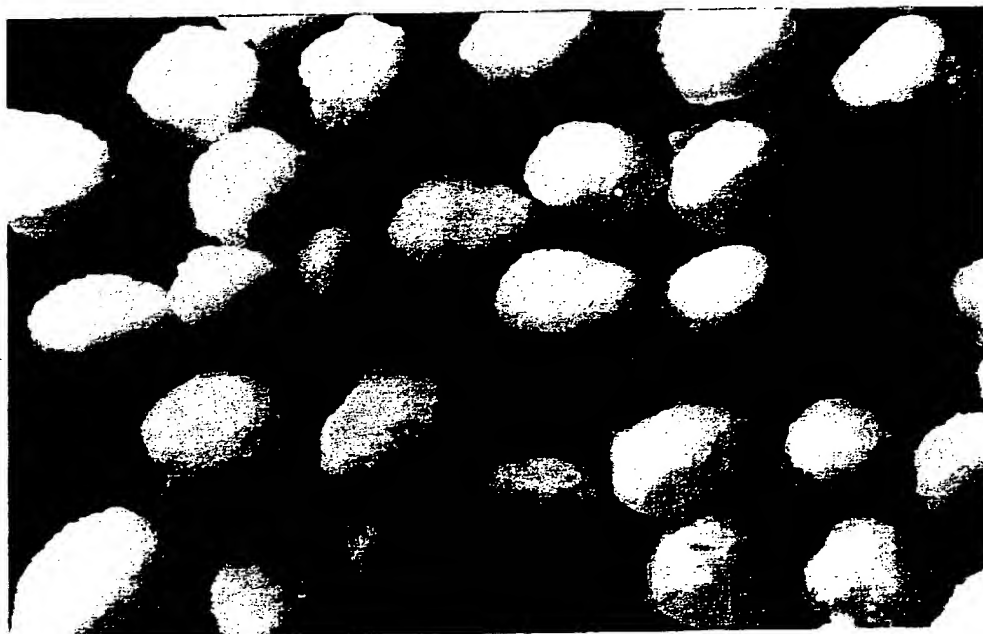
Fig.1

Microscopic Photos of Pyridoxine Spherical Granules

(A) Binder Solution: Water, X5.

(A)

1000 μ m



Microscopic Photos of Pyridoxine Spherical Granules

(B) Binder Solution: 70% Ethanol, X5.

(B)

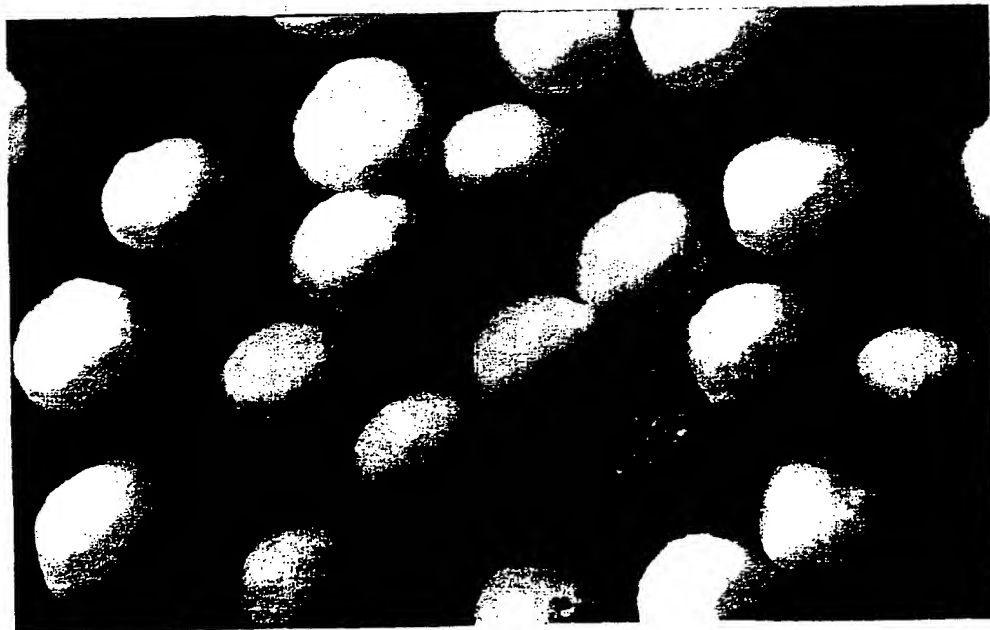


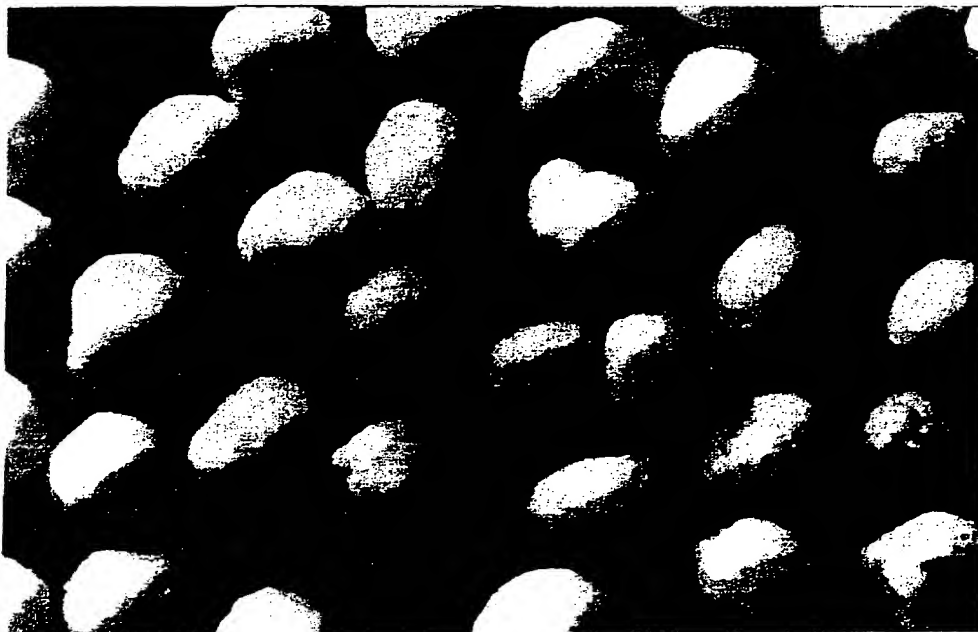
Fig.2

Microscopic Photos of Pyridoxine Spherical Granules

(A) Concentration of Microcrystalline Cellulose: 20%, X5.

(A)

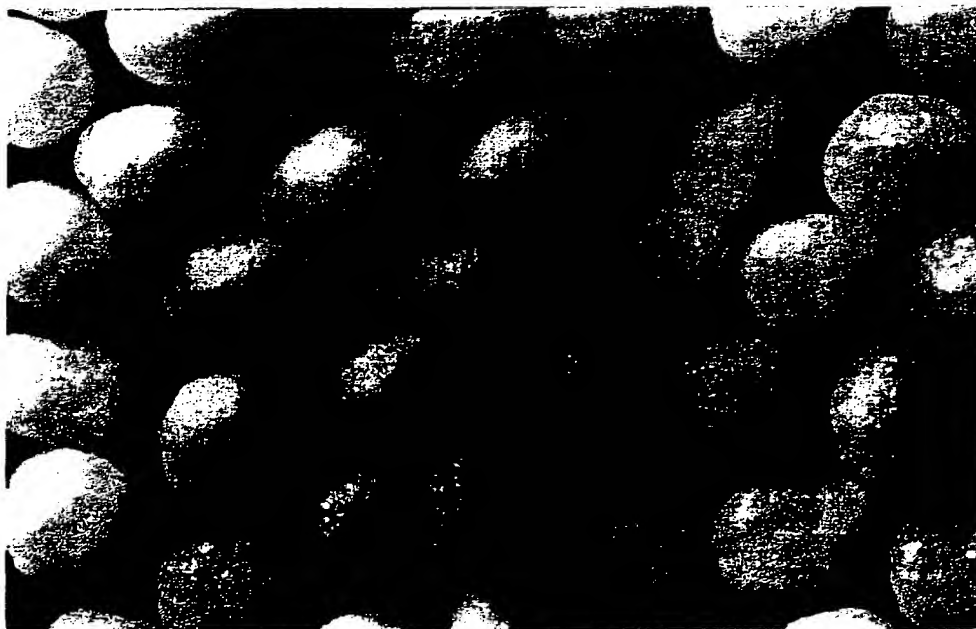
1000 μ m



Microscopic Photos of Pyridoxine Spherical Granules

(B) Concentration of Microcrystalline Cellulose: 75%, X5.

(B)



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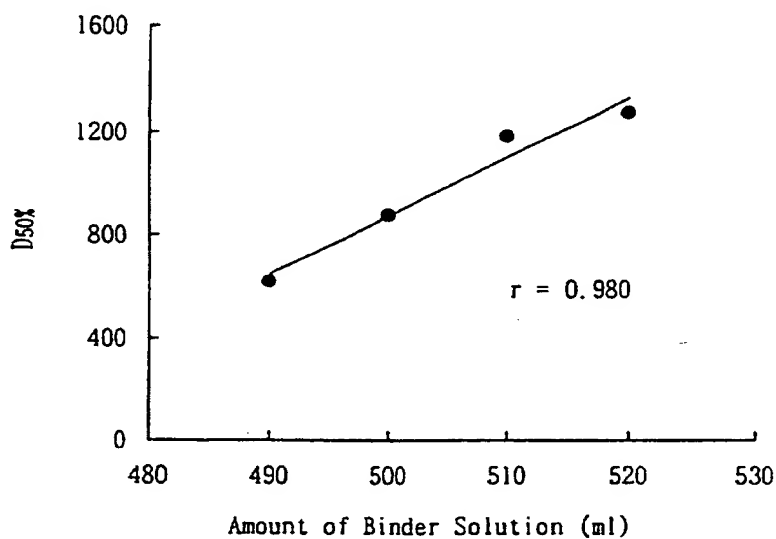


Fig. 3. Relationship between Amount of Binder Solution and D50% of Pyridoxine Spherical Granules

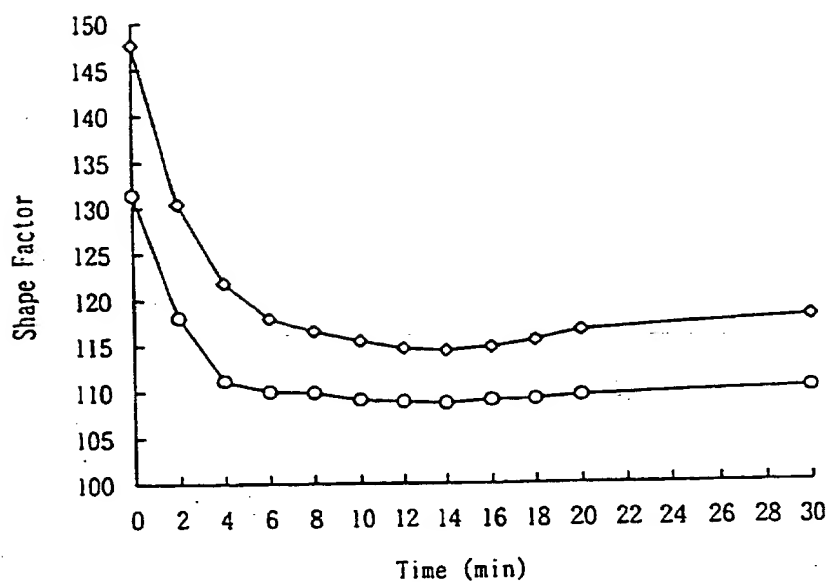


Fig. 4. Relationship between Agitation Time of Blade/Cross screw and Shape Factor of Pyridoxine Spherical Granules
◇, SF1; ○, SF2.

Fig.5

Microscopic Photos of Pyridoxine Spherical Granules

(A) Stirring Time: 0 Min, X5.

(A)

1000 μ m



Microscopic Photos of Pyridoxine Spherical Granules

(B-1) Stirring Time: 14 Min, X5.

(B-1)

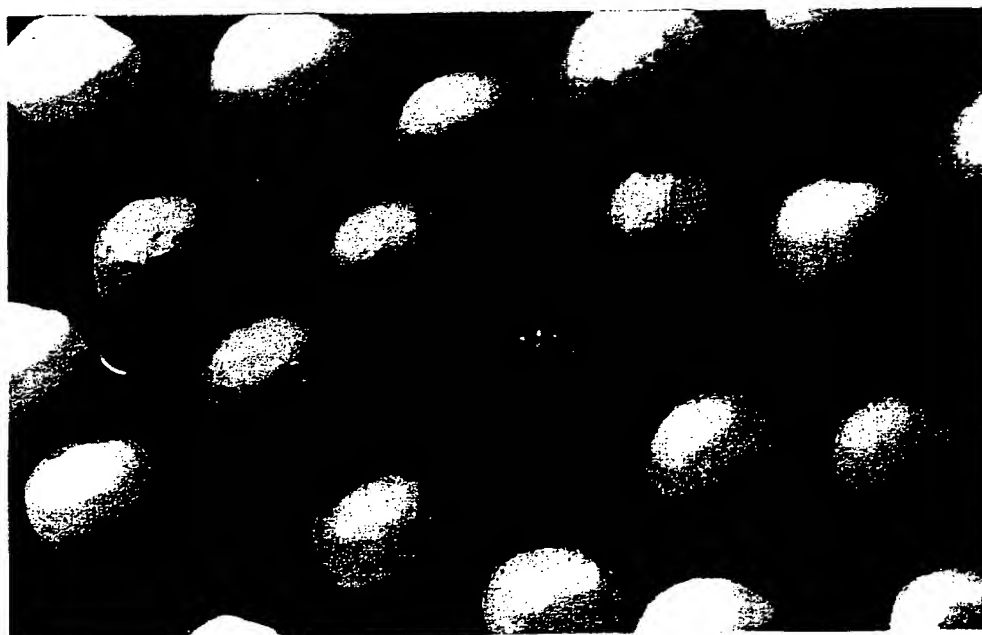
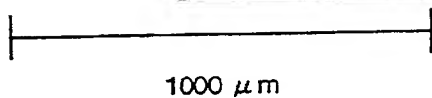
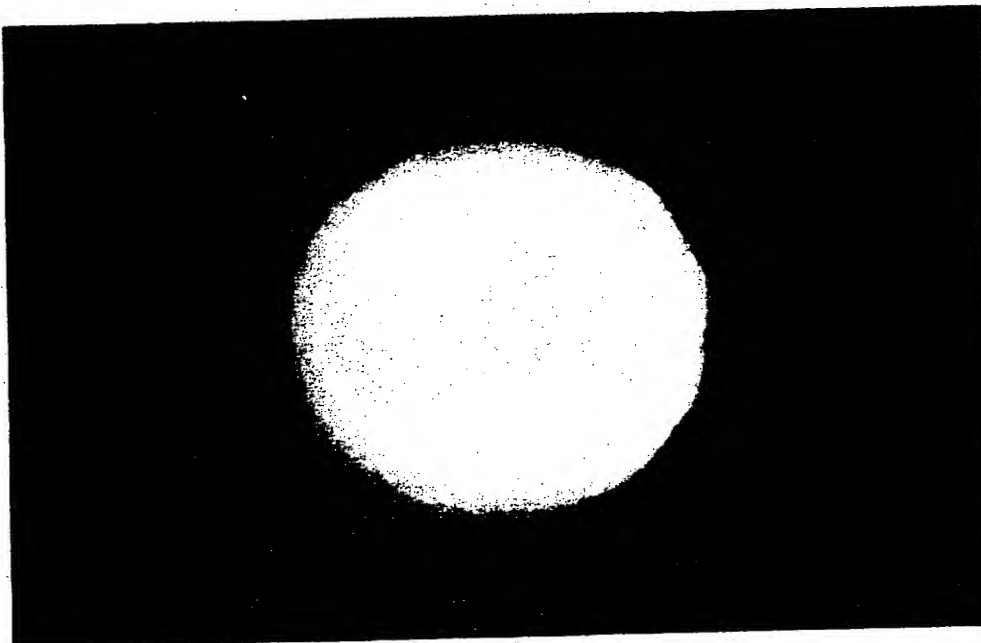


Fig.5

Microscopic Photos of Pyridoxine Spherical Granules

(B-2) Stirring Time: 14 Min, X15.

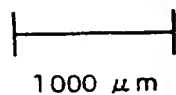
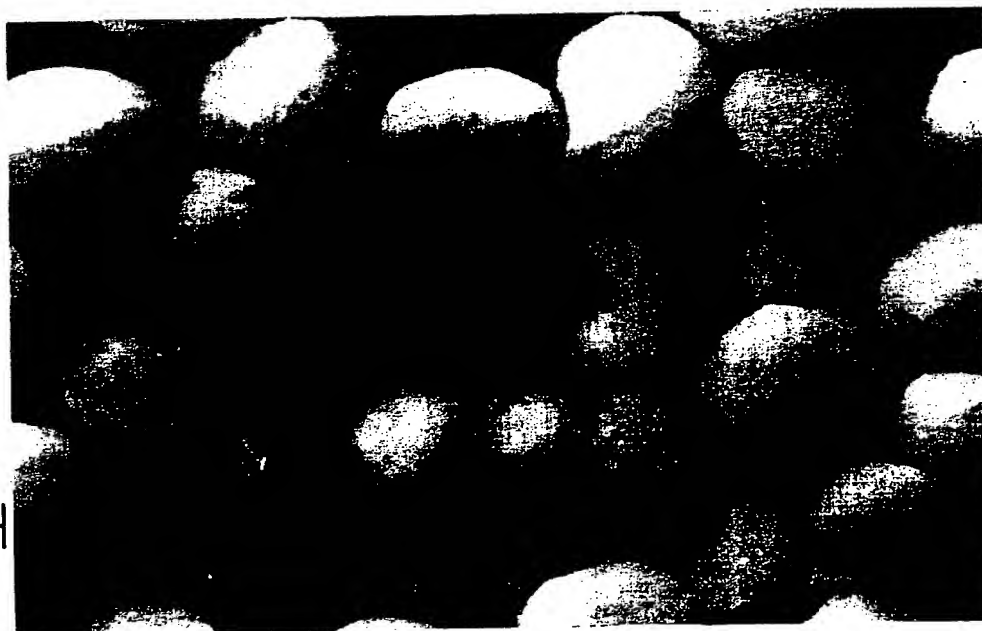
(B-2)



Microscopic Photos of Pyridoxine Spherical Granules

(C) Stirring Time: 30 Min, X5.

(C)



6/7

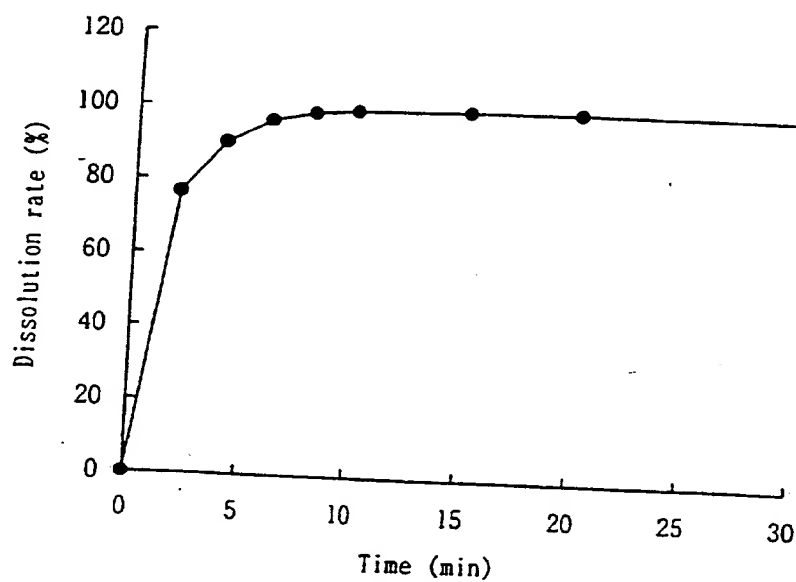
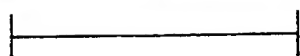


Fig. 6. Dissolution Curve of Pyridoxine Spherical Granules

Fig.7

Microscopic Photos of Pyridoxine Spherical Granules
after Dissolution Test, X10



1000 μ m



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INTERNATIONAL SEARCH REPORT

Int'l Application No
PCT/JP 96/02504

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 A61K9/16

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	EP 0 608 850 A (RECORDATI) 3 August 1994 see page 3, line 46 - line 55 see page 4, line 18 - page 5, line 7 see page 5, line 23 - line 24 ---	1,2,4-8
Y	EUROPEAN JOURNAL OF PHARMACEUTICS AND BIOPHARMACEUTICS, vol. 40, no. 1, February 1994, STUTTGART, DE, pages 32-35, XP000420350 JAN VERTOMMEN ET AL.: "Production of Pseudoephedrine HCl Pellets in a High Shear Mixer Granulator" see the whole document -----	1,2,4-8

☐ Further documents are listed in the continuation of box C.

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INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/JP 96/02504

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 608850 A	03-08-94	IT 1264020 B	09-09-96
		JP 6319786 A	22-11-94
		US 5460828 A	24-10-95

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